



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5 : A61K 31/725, 31/715, 31/70 C07H 1/00		A1	(11) International Publication Number: WO 92/20349 (43) International Publication Date: 26 November 1992 (26.11.92)
(21) International Application Number:	PCT/US92/04212	(74) Agent:	CLARK, Paul, T.; Fish & Richardson, 225 Franklin Street, Boston, MA 02110 (US).
(22) International Filing Date:	19 May 1992 (19.05.92)	(81) Designated States:	AT (European patent), AU, BE (European patent), CA, CH (European patent), DE (European patent), DK (European patent), ES (European patent), FI, FR (European patent), GB (European patent), GR (European patent), IT (European patent), JP, LU (European patent), MC (European patent), NL (European patent), NO, SE (European patent).
(30) Priority data:	703,254 20 May 1991 (20.05.91) US 833,973 11 February 1992 (11.02.92) US	Published	<i>With international search report.</i>
(71) Applicant:	GENZYME CORPORATION [US/US]; One Kendall Square, Cambridge, MA 02139 (US).		
(72) Inventors:	MILLER, Robert ; 24 Easterly Road, East Sandwich, MA 02537 (US). BURNS, James, W. ; 1630 Worcester Road, Apt. 410C, Framingham, MA 01740 (US). XU, Xuejian ; 402 Rindge Avenue 7M, Cambridge, MA 02140 (US).		

(54) Title: WATER INSOLUBLE DERIVATIVES OF POLYANIONIC POLYSACCHARIDES**(57) Abstract**

A water insoluble, biocompatible composition that is formed by a method which combines, in an aqueous mixture, a polyanionic polysaccharide and an activating agent, under conditions sufficient to form the composition; a nucleophile may also be added. Also, a water insoluble, biocompatible composition that is formed by a method which combines, in an aqueous mixture, a polyanionic polysaccharide and an activating agent under conditions sufficient to form the composition; a modifying compound and a nucleophile may also be added.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	FI	Finland	ML	Mali
AU	Australia	FR	France	MN	Mongolia
BB	Barbados	GA	Gabon	MR	Mauritania
BE	Belgium	GB	United Kingdom	MW	Malawi
BF	Burkina Faso	GN	Guinea	NL	Netherlands
BG	Bulgaria	GR	Greece	NO	Norway
BJ	Benin	HU	Hungary	PL	Poland
BR	Brazil	IE	Ireland	RO	Romania
CA	Canada	IT	Italy	RU	Russian Federation
CF	Central African Republic	JP	Japan	SD	Sudan
CG	Congo	KP	Democratic People's Republic of Korea	SE	Sweden
CH	Switzerland	KR	Republic of Korea	SN	Senegal
CI	Côte d'Ivoire	LI	Liechtenstein	SU	Soviet Union
CM	Cameroon	LK	Sri Lanka	TD	Chad
CS	Czechoslovakia	LU	Luxembourg	TG	Togo
DE	Germany	MC	Monaco	US	United States of America
DK	Denmark	MG	Madagascar		

- 1 -

WATER INSOLUBLE DERIVATIVES OF POLYANIONIC POLYSACCHARIDES

Background of the Invention

The present invention relates to biocompatible films
5 and gels formed from chemically modified polyanionic
polysaccharides.

Hyaluronic acid ("HA") is a naturally occurring
mucopolysaccharide found, for example, in synovial fluid, in
vitreous humor, in blood vessel walls and umbilical cord,
10 and in other connective tissues. The polysaccharide
consists of alternating N-acetyl-D-glucosamine and D-
glucuronic acid residues joined by alternating β 1-3
glucuronidic and β 1-4 glucosaminidic bonds, so that the
repeating unit is $-(1\rightarrow 4)-\beta-D-GlcA-(1\rightarrow 3)-\beta-D-GlcNAc-$. In
15 water, hyaluronic acid dissolves to form a highly viscous
fluid. The molecular weight of hyaluronic acid isolated
from natural sources generally falls within the range of
 5×10^4 up to 1×10^7 daltons.

As used herein the term "HA" means hyaluronic acid
20 and any of its hyaluronate salts, including, for example,
sodium hyaluronate (the sodium salt), potassium hyaluronate,
magnesium hyaluronate, and calcium hyaluronate.

HA, in chemically modified ("derivatized") form, is
useful as a surgical aid, to prevent adhesions or accretions
25 of body tissues during the post-operation period. The
derivatized HA gel or film is injected or inserted into the
locus between the tissues that are to be kept separate to
inhibit their mutual adhesion. To be effective the gel must
remain in place and prevent tissue contact for a long enough
30 time so that when the gel finally disperses and the tissue

- 2 -

do come into contact, they will no longer have a tendency to adhere.

Chemically modified HA can also be useful for controlled release drug delivery. Balazs et al., 1986, 5 U.S. Patent No. 4,582,865, states that "cross-linked gels of HA can slow down the release of a low molecular weight substance dispersed therein but not covalently attached to the gel macromolecular matrix." R.V. Sparer et al., 1983, Chapter 6, pages 107-119, in T.J. Roseman et al., Controlled 10 Release Delivery Systems, Marcel Dekker, Inc., New York, describes sustained release of chloramphenicol covalently attached to hyaluronic acid via ester linkage, either directly or in an ester complex including an alanine bridge as an intermediate linking group.

15 I. Danishefsky et al., 1971, Carbohydrate Res., Vol. 16, pages 199-205, describes modifying a mucopolysaccharide by converting the carboxyl groups of the mucopolysaccharide into substituted amides by reacting the mucopolysaccharide with an amino acid ester in the presence 20 of 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide hydrochloride ("EDC") in aqueous solution. They reacted glycine methyl ester with a variety of polysaccharides, including HA. The resulting products are water soluble; that is, they rapidly disperse in water or in an aqueous 25 environment such as is encountered between body tissues.

Proposals for rendering HA compositions less water soluble include cross-linking the HA. R.V. Sparer et al., 1983, Chapter 6, pages 107-119, in T.J. Roseman et al., Controlled Release Delivery Systems, Marcel Dekker, Inc., 30 New York, describe modifying HA by attaching cysteine residues to the HA via amide bonds and then cross-linking the cysteine-modified HA by forming disulfide bonds between the attached cysteine residues. The cysteine-modified HA

- 3 -

was itself water soluble and became water insoluble only upon cross-linking by oxidation to the disulfide form.

De Belder et al., PCT Publication No. WO 86/00912, describe a slowly-degradable gel, for preventing tissue adhesions following surgery, prepared by cross-linking a carboxyl-containing polysaccharide with a bi- or polyfunctional epoxide. Other reactive bi- or polyfunctional reagents that have been proposed for preparing cross-linked gels of HA having reduced water solubility include: 1,2,3,4-diepoxybutane in alkaline medium at 50°C (T.C. Laurent et al., 1964, *Acta Chem. Scand.*, vol. 18, page 274); divinyl sulfone in alkaline medium (E.A. Balasz et al., U.S. Patent No. 4,582,865, (1986); and a variety of other reagents including formaldehyde, dimethylolurea, dimethylolethylene urea, ethylene oxide, a polyaziridine, and a polyisocyanate (E.A. Balasz et al., U.K. Patent Appl. No. 84 20 560 (1984). T. Mälson et al., 1986, PCT Publication No. WO 86/00079, describe preparing cross-linked gels of HA for use as a vitreous humor substitute by reacting HA with a bi- or polyfunctional cross-linking reagent such as a di- or polyfunctional epoxide. T. Mälson et al., 1986, EPO 0 193 510, describe preparing a shaped article by vacuum-drying or compressing a cross-linked HA gel.

25 Summary of the Invention

In one aspect, the invention features a method for making a water insoluble biocompatible composition, the method including combining, in an aqueous mixture, a polyanionic polysaccharide, an activating agent, and a nucleophile, under conditions sufficient to form the composition.

In preferred embodiments of this aspect of the invention, the activating agent which are used include

- 4 -

benzotriazole-1-yloxytris (dimethylamino)phosphonium hexafluorophosphate, O-benzotriazole-1-yl-N,N,N',N'-tetramethyluronium hexafluorophosphate, bromo-tris(dimethylamino)phosphonium hexafluorophosphate, or the 5 corresponding halide salts thereof.

The preferred concentration of polyanionic polysaccharide in the reaction is 0.0002 - 0.1M, more preferably 0.0005 - 0.02M. The preferred pH for carrying out the reaction is 3.5 to 8.0. The preferred reagent 10 stoichiometry is at least 0.1 molar equivalents of activating agent per molar equivalent of polyanionic polysaccharide.

Another aspect of the invention features a method for making a water insoluble biocompatible composition, the 15 method including combining, in an aqueous mixture, a polyanionic polysaccharide, an activating agent, a modifying compound, and a nucleophile, under conditions sufficient to form the composition.

In preferred embodiments of this aspect of the 20 invention, modifying compounds include, 1-hydroxy-benzotriazole hydrate, 1-hydroxybenzotriazole monohydrate, N-hydroxysulfosuccinimide, N-hydroxysuccinimide, 4-nitrophenol, 2-nitrophenol, 4-nitrothiophenol, 25 2-nitrothiophenol, pentachlorophenol, pentafluorophenol, imidazole, tetrazole, 4-dimethylaminopyridine or other related compounds. The activating agent is preferably a diimide, more preferably a carbodiimide, e.g., 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide and 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide methiodide.

30 Also in preferred embodiments of the second aspect of the invention, the preferred polyanionic polysaccharide is present in the reaction at a concentration of 0.0002 - 0.1M, more preferably 0.005 - 0.02M. The preferred pH for

- 5 -

carrying out the reaction is 3.5 to 8.0. The preferred reagent stoichiometry is at least 0.1 molar equivalents of activating agent per molar equivalent of polyanionic polysaccharide, and at least 1 molar equivalent of modifying compound per molar equivalent of activating agent.

Preferred polyanionic polysaccharides for use in the methods of the invention include hyaluronic acid (HA), carboxymethyl cellulose (CMC), carboxymethyl amylose (CMA), chondroitin-6-sulfate, dermatin sulfate, heparin, and heparin sulfate; HA, CMC, and CMA are particularly preferred. It is also well understood that two or more polyanionic polysaccharides may be employed in the methods of the invention.

Also in both aspects of the invention, preferred nucleophilic compounds which are capable of reacting with the activated polyanionic polysaccharide include amino acid amides (preferably leucinamide hydrochloride), monofunctional amines (preferably 3-amino-1-propanol), amino acid esters (preferably a methyl ester or a butyl ester, including t-butyl ester), amino alcohols, amino thiols, amino phenols, amino cathechols, amino acids, salts of amino acids, peptides, proteins and other ambident nucleophilic compounds in which only one electron rich moiety reacts as a nucleophile with the activated polyanionic polysaccharide.

In yet another aspect, the invention features a method for preparing a water insoluble composition by combining a polyanionic polysaccharide and an activating agent under conditions sufficient to form the composition.

Preferred polyanionic polysaccharides according to this aspect of the invention include hyaluronic acid (HA), carboxymethylcellulose (CMC), carboxymethylamylose (CMA), chondroitin-6-sulfate, dermatin sulfate, heparin, and heparin sulfate; HA, CMC, and CMA are particularly preferred.

- 6 -

The preferred pH for carrying out the reaction is 3.5 to 8.0, more preferably 4.0 to 5.1 and most preferably 4.7 to 5.1. The preferred concentration for the polysaccharide is 0.005 - 0.1M, more preferably 0.01 - 5 0.02M. The molar ratio of polysaccharide to activating agent is preferably at least 1:1, more preferably about 1:4. The preferred activating agent is a carbodiimide, e.g., 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide and 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide methiodide.

10 The term "aqueous mixture", as used herein, generally refers to a solution composed primarily of water, but which may also comprise as much as 1 part in 20 of a polar aprotic solvent. Preferred aprotic solvents include acetonitrile, dimethylformamide, hexamethylphosphoramide, 15 dimethylacetamide, *N*-methylpyrrolidinone, 1,4-dioxane, and acetone.

A "polyanionic polysaccharide" is a polysaccharide containing more than one negatively charged group, e.g., carboxyl groups at pH values above about pH 4.0.

20 The terms "mole or molar concentration(M)" of polyanionic polysaccharides, as used herein, refer to moles of the repeating monomeric unit contained within the polymer.

A polyanionic polysaccharide is said to be 25 "activated", as that term is used herein, when it is treated in an aqueous mixture in a manner that renders the carboxyl groups on the polyanionic polysaccharide vulnerable to nucleophilic attack; and an "activating agent" is a substance that, in an aqueous mixture including a 30 polyanionic polysaccharide, causes the polyanionic polysaccharide to become so activated.

A "modifying" compound is defined as a reagent which, in the presence of an activated polyanionic

- 7 -

polysaccharide, reacts with the activated carboxyl moiety of the polyanionic polysaccharide to form a new activated species capable of reacting with a nucleophile.

The activated polyanionic polysaccharides which 5 comprise the water insoluble compositions produced by the method of the invention may be in the form of a gel, or in the form of fibers. Blends may also be prepared by mixing various amounts of two or more different activated-polyanionic polysaccharides. Preferably, blends consist of 10 activated-HA and activated-CMC, or activated-HA and activated-CMA.

The compositions and blends of the invention may be provided in the form of an adhesion prevention composition, e.g., in the form of a film, foam, or composition suitable 15 for incorporation in a syringe. They may also include a pharmaceutically active substance dispersed throughout making them useful as a drug delivery system. Suitable substances include proteins, growth factors, enzymes, drugs, biopolymers, and biologically compatible synthetic polymers.

20 The term "film", as used herein, means a substance formed by compressing a gel or fibers, or by allowing or causing a gel or fibers to dehydrate. Any gel or fibers of the invention may be formed into such a film.

The term "foam", as used herein, means a substance 25 formed by introducing gas bubbles into the gels or fibers of the invention.

A "biocompatible" substance, as the term is used herein, is one that has no medically unacceptable toxic or injurious effects on biological function.

30 We have discovered that a gel, foam, or film produced by treating a polyanionic polysaccharide with a suitable activating agent, may be made having decreased

- 8 -

water solubility, without the use of any separately added bi- or polyfunctional cross-linking agent.

A "water soluble" gel, or film, as that term is used herein, is one which, formed by drying an aqueous 5 solution of 1% weight/weight ("w/w") sodium hyaluronate in water, having dimensions 3 cm x 3 cm x 0.3 mm, when placed in a beaker of 50 ml of distilled water at 20°C. and allowed to stand without stirring, loses its structural integrity as a film after 3 minutes, and becomes totally dispersed within 10 minutes. A "water insoluble" film of the invention, as that phrase and like terms are used herein, formed using a 1% aqueous solution of a polyanionic polysaccharide, modified according to the invention, having the same 15 dimensions and similarly allowed to stand without stirring in a beaker of 50 ml of distilled water at 20°C., is structurally intact after 20 minutes; the film boundaries and edges are still present after 24 hours, although the film is swollen.

Because the gels and films are water insoluble, 20 they can be thoroughly washed with water before use to remove unreacted substances.

Gels, foams, and films of the invention can also be prepared in colored form, by including a dye or stain in the reaction mixture. Such colored films and gels can be more 25 easily seen when in place or during placement, making them easier to handle during surgical procedures than colorless ones.

The films, gels, and foams of the invention retain their strength even when hydrated. Because they adhere to 30 biological tissues without the need for sutures, they are useful as postoperative adhesion prevention membranes. They can be applied to tissue even in the presence of bleeding.

- 9 -

Other features and advantages of the invention will be apparent from the following description of the preferred embodiments thereof, and from the claims.

Description of the Preferred Embodiments

5 Lysine-Modified HA

The gels, foams, and films of the invention are made generally as follows. HA is dissolved in water and the pH of the resulting aqueous mixture is adjusted downward; then the dissolved HA is activated by admixing a suitable 10 activating agent, and a suitable lysine ester is admixed with the activated HA and allowed to stand until the desired gel has formed. The activating agent and the ester can be admixed in any sequence.

15 The preferred method of making the lysine-modified gels and films of the invention will now be described in more detail. As one skilled in the art will appreciate, gels and films of the invention can be made using protocols that are within the method of the invention yet are different in particulars from those described here.

20 A sample of hyaluronic acid or a salt of hyaluronic acid, such as sodium hyaluronate, is dissolved in water to make an aqueous mixture. HA from any of a variety of sources can be used. As is well-known, HA can be extracted from animal tissues or harvested as a product of bacterial 25 fermentation. Hyaluronic acid can be produced in commercial quantities by bioprocess technology, as described for example in PCT Publication No. WO 86/04355. Preferably the concentration of HA in this first aqueous mixture is in the range between 0.4% and 2.5% weight/weight ("w/w").
30 Subsequent reactions are slower and less effective at significantly lower concentrations, while significantly higher concentrations are difficult to handle owing to their high viscosity.

- 10 -

The aqueous HA mixture should be acidic, preferably having a pH between pH 4.0 and pH 5.0, more preferably between pH 4.3 and pH 4.75. At lower pH values the preferred activating agent, EDC, is unstable, and at higher 5 values the reaction rate is diminished. Preferably hydrochloric acid is added to adjust the pH, although other known acids can be used.

Once the pH of the aqueous HA mixture has been adjusted, an activating agent is admixed. Preferred 10 activating agents include carbodiimides, most preferably EDC (in some references this substance is termed 1-(3-dimethylaminopropyl)-3-ethyl-carbodiimide or "DEC") or ETC (1-ethyl-3-(3-dimethylaminopropyl)carbodiimide methiodide).

Then a nucleophilic lysine ester is admixed to the 15 aqueous HA-activating agent mixture. Preferred esters include methyl, ethyl, or t-butyl esters. The lysine can be in the form of di-lysine, tri-lysine, or polylysine, or their hydrochloride salts.

The lysine ester and the activating agent may be 20 admixed to the pH adjusted HA mixture in any sequence, either all at once or gradually.

If a colored product is desired, a solution of a dye or stain such as the blue dye "Brilliant Blue R", also known as "Coomassie" Brilliant Blue R-250", distributed as "Serva 25 Blue" by Serva, can be admixed to the reaction mixture at this point. The resulting product has a blue color that can provide a good contrast to the color of body tissues, making the film or gel easy to see while it is handled during surgery and once it is in place.

Once the reagents (and the stain or dye, if any) 30 have been admixed, the reaction mixture can be simply allowed to stand for a time, or it can be continually or occasionally stirred or agitated.

- 11 -

Upon admixing of the reagents the pH rises, and can be maintained at the desired pH by addition of acid as the reaction proceeds. We have found, however, that films and gels with various desired physical properties can be obtained by simply allowing the pH to rise as the reaction proceeds. The mode of addition of the reagents, particularly the EDC and the lysine ester, is not critical, but the ratios of these reagents to the HA is important. We have found that the best results are obtained when the ratio of HA:EDC:Lysine ester ranges from 1:2:1 to 1:4:10. Lower values typically result in weaker, less insoluble products, while higher values typically result in stronger, more insoluble products.

Polyanionic Polysaccharide-Modified HA

15 Polyanionic polysaccharide-modified HA gels and films are prepared generally by mixing HA (as described above) with a polyanionic polysaccharide and an activating agent to form a water-insoluble precipitate. The precipitate can be cast into thin membranes useful for 20 postoperative adhesion prevention. It can also be colored as described above. To increase the strength of films cast from the precipitate, the films may be subjected to dehydrothermal treatment in which they are heated under vacuum (about 30 mm Hg) at approximately 105°C for 24 hr.

25 The polysaccharide and HA can be mixed together, after which the activating agent is added. Alternatively, the polysaccharide may be reacted with the activating agent, followed by addition of HA. A third option is to combine the HA with the activating agent, followed by addition of 30 the polysaccharide. Preferred activating agents are as described above and include the carbodiimides EDC and ETC. The reaction is preferably carried out at a pH between 4 and 5. The preferred polysaccharide concentration ranges from

- 12 -

0.005 to 0.1M, and is more preferably in the range 0.01 to 0.02M. The preferred molar ratio of polysaccharide to activating agent is at least 1:1, more preferably about 1:4.

Activated Polyanionic Polysaccharides

5 Polyanionic polysaccharide gels, films, and foams are prepared generally by mixing at least one polyanionic polysaccharide (e.g., HA, CMC, CMA) with an activating agent to form a water-insoluble material. Preferred activating agents include the carbodiimides, EDC and ETC. The reaction 10 may be carried out at a pH between 3.5 and 8, with optimal reaction conditions occurring between pH 4.7 and 5.1. The polysaccharide molecular weight used in the reaction may range from 9.0×10^4 to 3.0×10^6 daltons, but preferably is between 2.5×10^5 to 1.0×10^6 daltons. The preferred molar 15 ratio of polysaccharide to activating agent is at least 1:1, and more preferably about 1:4. The insoluble material formed by this method may be in the form of a gel or in the form of fibers and can be used directly for adhesion prevention or drug delivery, or can be cast onto flat molds 20 and either air dried or lyophilized to yield thin films or foams.

In addition, blends can be prepared by mixing various amounts of different unpurified or purified activated-polyanionic polysaccharides. These blends are 25 made homogeneous by mixing with overhead stirrers and/or high shear mixers. Unreacted activating agent may be removed from the unpurified mixture by molecular weight sizing, dialysis, dialfiltration or fractional precipitation with a water-soluble solvent, according to standard methods, 30 prior to use. The purified mixture can be used directly for adhesion prevention and/or drug delivery, or may be cast

- 13 -

onto flat molds and either air dried or lyophilized to form films or foams.

Bop reagent-activated Polyanionic Polysaccharides

5 Polyanionic polysaccharide water-insoluble gels, films, and foams are also prepared generally by dissolving at least one polyanionic polysaccharide (e.g. HA, CMC, CMA) in an aqueous mixture; activating the polyanionic polysaccharide with an activating agent such as benzotriazole-1-yloxytris(dimethylamino)phosphonium 10 hexafluorophosphate (Bop-reagent); and reacting the activated polyanionic polysaccharide with a suitable nucleophile to form the desired insoluble composition.

15 The reaction may be carried out at a pH between 3.5 and 8, with optimal reaction conditions between pH 4.6 and 5.0. The molecular weight of the polyanionic polysaccharide used in the reaction may range from 6.0×10^2 to 4×10^6 daltons, but preferably greater than 5×10^5 daltons. The preferred reagent stoichiometry is at least 0.1 molar equivalents of activating agent per molar equivalent of 20 polyanionic polysaccharide, and at least 1 molar equivalent of nucleophile per molar equivalent of polyanionic polysaccharide.

25 One major unexpected advantage of the BOP activation of polyanionic polysaccharide is that the molecular weight of the polyanionic polysaccharide is not decreased upon coupling to the nucleophile. This result is in contrast to reactions involving carbodiimides solely, in which we observe a decrease in HA molecular weight upon nucleophilic coupling.

30 A non-obvious aspect of this invention is that organic soluble activating agents, described previously for use only in organic solvents, are able to effect, in an

- 14 -

aqueous milieu, chemical coupling of a nucleophile with the water-soluble polyanionic polysaccharides. This observation presents a significant additional unexpected advantage in that any unreacted activators can be removed from the water-
5 insoluble product simply by extracting the reaction solution with any appropriate water-immiscible, organic solvent. Examples of such solvents may include diethyl ether, methylene chloride, chloroform, ethyl acetate, or tetrahydrofuran.

10 Modified Carbodiimide-activated Polyanionic Polysaccharides

Polyanionic polysaccharide water-insoluble gels, films, and foams are prepared generally by dissolving at least one polyanionic polysaccharide (e.g., HA, CMC, CMA), in an aqueous mixture; activating the polyanionic
15 polysaccharide with an activating agent such as a diimide, e.g. EDC or ETC; modifying the activated polyanionic polysaccharide with a modifying compound such as 1-hydroxybenzotriazole hydrate (HOBr), 1-hydroxybenzotriazole monohydrate or one of the other
20 compounds described above; and reacting the activated polyanionic polysaccharide with a suitable nucleophile to form the desired insoluble composition.

The molecular weight range of the polyanionic polysaccharide used in the reaction may be 6.0×10^2 to 4.0×10^6 daltons, but preferably is greater than 5.0×10^5 daltons. The preferred reagent stoichiometry is at least 0.1 molar equivalents of a activating compound per molar equivalent of polyanionic polysaccharide, and more preferably at least 1:1. The preferred reagent
30 stoichiometry also encompasses at least one molar equivalent of modifying compound per molar equivalent of activating agent. The reaction with the activating agent may be

- 15 -

carried out at a pH between 3.5 and 8, with optimal reaction conditions occurring between 4.0 and 4.8. The modification of the polyanionic polysaccharide with the modifying compound is carried out at a pH between 3.0 and 8.0, with 5 optimal modification conditions occurring between pH 3.3 and 4.5.

In the case where the activating agent used is a diimide, the modifying compound reacts with the intermediate O-acylisourea to form a new active carbonyl group capable of 10 being transferred to a nucleophile.

It is our discovery that organic soluble modifying reagents, which display low solubility in water, can effect the coupling of polyanionic polysaccharides to nucleophiles in an aqueous mixture. A significant unexpected advantage 15 of this solubility difference between modifier and polyanionic polysaccharide is that the insoluble product can be purified by extraction with a suitable, water-immiscible organic solvent such as chloroform, methylene chloride, ethyl acetate, or diethyl ether, or by alcohol precipitation 20 and trituration.

An additional unexpected advantage imparted by a coupling of nucleophiles to a modified, activated polyanionic polysaccharide is that the reaction can be carried out under acidic conditions which provides control 25 of the specificity in reactions with ambident nucleophilic compounds (e.g., amino alcohols, amino acids, amino thiols, amino phenols, amino catechols, peptides, and proteins) so that only one of potentially several electron rich moieties are capable of reacting as a nucleophile with the activated 30 polyanionic polysaccharide.

Films and Gels

Polyanionic polysaccharides modified according to the above descriptions can be cast as films in a

- 16 -

straightforward manner. Typically the reaction mixture is poured into a vessel having the desired size and shape and allowed to air dry. In general films formed by drying mixtures poured thickly, so that they have a lower surface area/volume, possess greater strength than films formed by drying thinner, higher surface area/volume mixtures.

Alternatively a film can be formed by compressing a gel under conditions that permit escape of water, as, for example, by compressing the gel between two surfaces, at least one of which is porous, as described, for example, in EPO 0 193 510.

If desired, a gel or film can be washed prior to use by, for example, perfusion with water or 1 M aqueous sodium chloride. Alternatively the reaction mixture can be dialyzed to remove residual reagents prior to casting as a film. Washing to remove residual reagents or reagent-derived material such as substituted ureas is desirable if the film or gel is to be used for therapeutic applications. Gels or films colored blue with Brilliant Blue R as described above do not lose their coloration during such washing. The removal of reagents or reaction products can be monitored by high pressure liquid chromatography.

Detailed Description of the Invention

The invention is described in more detail in the following examples. These examples are given by way of illustration and are not intended to limit the invention except as set forth in the claims.

Example 1: In this example hydrogels were prepared using EDC as an activating agent and leucine methyl ester hydrochloride as a nucleophile.

Sodium hyaluronate (400 mg; 1.0 mmol of carboxyl groups) having a molecular weight between 1×10^6 and $2 \times$

- 17 -

10^6 was dissolved in 10 ml of distilled water. The pH of the aqueous solution was adjusted to pH 4.75 by addition of 0.1 N HCl. Then 314 mg of EDC (1.64 mmol) was added all at once followed by 190 mg (1.05 mmol) of L-leucine methyl ester hydrochloride. The pH of the reaction mixture then rose to 6.2 over two hours. The reaction mixture was kept at room temperature for five hours, after which time it had formed a thick insoluble hydrogel. This hydrogel could be washed with a 1 M NaCl solution to remove residual reagents without loss of its physical properties.

Example 2: In this example various EDC/leucine:HA ratios were used for comparison of gel formation and properties.

The procedure was as in Example 1, using sodium hyaluronate (400 mg; 1.0 mmol of carboxyl groups) in 15 ml of water. In separate experiments the following quantities of EDC and leucine methyl ester hydrochloride were then added: 153 mg EDC (0.8 mmol)/182 mg leucine methyl ester hydrochloride (1.0 mmol); 76 mg EDC (0.4 mmol)/90 mg leucine methyl ester hydrochloride (0.5 mmol); and 38 mg EDC (0.2 mmol)/45 mg leucine methyl ester hydrochloride (0.25 mmol). Strong hydrogels were obtained as in example 1 for the highest ratio of EDC and leucine methyl ester hydrochloride. At the lowest ratio of reactants (0.2 mmol/0.25 mmol to 1.0 mmol HA carboxyl groups) a weak gel was obtained, which collapsed to a fluid after two weeks.

Example 3: In this example the HA concentration was reduced by one-half for comparison of resulting gel properties.

The procedure was as in example 1 except the HA (400 mg; 1.0 mmol of carboxyl groups) was dissolved in 30 ml of water rather than 15 ml (1-1/3% w/w HA). A hydrogel was

- 18 -

formed, although it was weaker than that obtained in Example 1.

Example 4: In this example films were prepared using EDC as an activating agent and leucine methyl ester hydrochloride as a nucleophile.

Sodium hyaluronate (400 mg; 1.0 mmol of carboxyl groups) was dissolved in 40 ml of distilled water. The pH of the solution was adjusted to pH 4.75 by addition of 0.1 N HCl. Then EDC (314 mg; 1.64 mmol) was added in a single portion, followed by 190 mg (1.05 mmol) of L-leucine methyl ester hydrochloride. The pH of the reaction mixture rose to 6.2 during two hours, after which time the solution was poured into a petri dish of area 6360 mm², and allowed to dry to a film over a two day period. Films produced in this manner were strong and insoluble in water and 1 M aqueous NaCl. The films could be washed with water or aqueous NaCl as in Example 1 to remove residual reagents without loss of their physical properties. Infrared spectroscopic analysis of such films showed no carbodiimide absorption at about 2130 cm⁻¹ and displayed absorptions at about 1740 cm⁻¹, 1700 cm⁻¹, 1650 cm⁻¹, and 1550 cm⁻¹.

Example 5: In this example various HA concentrations were used in making films for comparison of resulting film properties.

The procedure described in example 4 was repeated, using three different initial HA concentrations made by dissolving the HA (400 mg; 1.0 mmol of carboxyl groups) in 30 ml, 40 ml, or 100 ml of distilled water. Films produced using each of these initial concentrations of HA were strong and insoluble in water and 1 M aqueous NaCl, showing that a range of concentrations of HA can be used. Each of these

- 19 -

films could be washed with water or aqueous NaCl without loss of its physical properties.

Example 6: This example illustrates the effect of dialyzing the reaction mixture prior to casting to form a 5 film, as compared with washing the film after forming it.

Sodium hyaluronate (400 mg in 40 ml of water), EDC (314 mg; 1.64 mmol) and L-leucine methyl ester hydrochloride (190 mg; 1.05 mmol) were allowed to react as in Example 4. Upon completion of reaction (2 hours) the reaction mixture 10 was dialyzed against water, through 12,000 NMW cutoff dialysis tubing in order to remove residual reagents. The dialyzed mixture was then cast as a film as in Example 4. The film so obtained was strong and insoluble in water or 1 M aqueous NaCl.

15 Example 7: In this example films were formed by drying more thickly poured reaction mixtures, to compare the properties of films produced from drying mixtures at differing surface area/volume.

A reaction mixture obtained as in Example 4 (40 ml 20 reaction volume) was cast into a small petri dish (area 3330 mm²). The film so obtained was insoluble in 1 M aqueous NaCl and in water (100°C; 1 hour).

Example 8: In this example films were prepared using other amino acid esters and HA activated with EDC.

25 A solution of HA (400 mg in 40 ml of H₂O) was brought to pH 4.7 using 0.1 N HCl. Then EDC (314 mg; 1.6 mmol) was added all at once followed by 1 mmol of the amino acid derivative. The reaction mixture was poured into a petri dish and allowed to dry. Insoluble films were 30 obtained from L-valine methyl ester hydrochloride, L-isoleucine methyl ester hydrochloride, L-proline methyl

- 20 -

ester hydrochloride, and L-phenylalanine methyl ester hydrochloride.

Example 9: In this example films were prepared using a simple primary amine (aniline) as a nucleophile.

5 A solution of HA (400 mg in 40 ml of H₂O) was brought to pH 4.7 using 0.1 N HCl. Then EDC (314 mg; 1.6 mmol) was added all at once followed by 1 mmol of aniline. The reaction mixture was poured into a petri dish and allowed to dry, and insoluble films were obtained.

10 Example 10: In this example films were prepared using other esters of leucine.

A solution of HA (400 mg in 40 ml of H₂O) was brought to pH 4.7 using 0.1 N HCl. Then EDC (314 mg; 1.6 mmol) was added all at once followed by 1 mmol of the 15 leucine ester. The reaction mixture was poured into a petri dish and allowed to dry. Insoluble films were obtained from both L-leucine ethyl ester hydrochloride and L-leucine t-butyl ester hydrochloride.

Example 11: In this example gels were prepared 20 using other amino acid methyl esters.

A solution of HA (400 mg in 15 ml of H₂O) was brought to pH 4.7 and EDC (314 mg; 1.6 mmol) was added, followed by the amino acid derivative (1 mmol). The reaction mixture formed a thick gel within from 5 to 24 25 hours. Water insoluble gels were obtained using L-valine methyl ester hydrochloride, L-isoleucine methyl ester hydrochloride, L-arginine methyl ester hydrochloride, L-proline methyl ester hydrochloride, and L-histidine methyl ester hydrochloride.

Example 12: In this example films were prepared 30 using an amino acid amide (leucinamide) as a nucleophile.

- 21 -

A solution of HA (400 mg in 40 ml of H₂O) was brought to pH 4.7 using 0.1 N HCl. Then EDC (314 mg; 1.6 mmol) was added all at once followed by 1 mmol of L-leucinamide hydrochloride. The reaction mixture was poured 5 into a petri dish and allowed to dry. and insoluble films were obtained.

Example 13: In this example gels were prepared using leucine ethyl ester hydrochloride.

A solution of HA (400 mg in 15 ml of H₂O) was 10 brought to pH 4.7 and EDC (314 mg; 1.6 mmol) was added, followed by leucine ethyl ester hydrochloride (1.0 mmol). The mixture formed a thick, water insoluble gel within from 5 to 24 hours.

Example 14: In this example films and gels were 15 prepared using ETC as the HA activating agent.

Sodium hyaluronate (400 mg, 1.0 mmol of carboxyl groups) having a molecular weight in the range between 1 x 10⁶ and 2 x 10⁶ daltons was dissolved in water (10 ml and 30 ml). The pH of each aqueous solution was adjusted to pH 20 4.75 by addition of 0.1 N HCl. Then 475 mg of ETC (1.6 mmol) was added all at once, followed by 190 mg (1.05 mmol) of L-leucine methyl ester hydrochloride. The pH of this reaction mixture rose to pH 6.2 over the next 2 hours. The reaction mixture containing 10 ml of water formed an 25 insoluble gel. The reaction mixture containing 30 ml of water gave an insoluble film after drying.

Example 15: This example illustrates the preparation of a colored film.

A solution of HA (400 mg in 30 ml of H₂O) was 30 brought to pH 4.75 as in example 13 and then ETC (475 mg; 1.6 mmol) and leucine methyl ester hydrochloride (190 mg; 1.05 mmol) were added. A dilute solution of "Serva Blue" (5

- 22 -

mg/ml) dye in H₂O (0.5 ml) was then added to the reaction mixture. The resulting mixture was poured into a Petri dish and a water insoluble blue film was obtained after 16 hours. The blue color was retained by the film when the 5 film was washed with 1 M NaCl and then with H₂O.

Example 16: This example illustrates the tissue biocompatibility of a film of chemically modified HA.

Four strips of films prepared according to the procedure described in Example 4, and two USP negative 10 control strips were surgically implanted into the paravertebral muscle of White New Zealand rabbits (two per test). The test sites were evaluated either macroscopically after 72 hours or with complete histopathology after 7 days. In accordance with the USP XXI, p. 1237, the test 15 material met the requirements of the USP Implantation Test for the Evaluation of Plastic Materials.

Example 17: This example illustrates the preparation of lysine-modified HA.

A 0.4% (w/w) solution of HA in water was prepared. 20 The pH of this solution was adjusted to between 4.3 and 4.75 by addition of acid. To each 100 ml of this solution was added 0.76 g of EDC with stirring until the EDC had completely dissolved. To each 100 ml of the HA/EDC solution was added 0.20 g of lysine methyl ester (LME) with stirring 25 until the LME had completely dissolved. The addition of HA, EDC, and LME was conducted at room temperature; once the final HA/EDC/LME solution had been formed, it was stored at 4°C until needed.

The LME-modified HA material can be processed into 30 various shapes, sizes, and consistencies depending on the end application. If a thin sheet of the material is desired, the mixture can be poured onto a flat surface.

- 23 -

This material can then be turned into a solid by allowing the water to evaporate under ambient or elevated temperatures. An alternative method of producing sheets of the material is to subject it to freeze drying. The pore 5 size of the final product can be controlled by adjusting the initial freezing temperature. Curved surfaces and other shapes can be produced in a similar manner by initially casting the gel onto a negative image surface and then processing as described. The dried sheet can be processed 10 further, if desired, by pressing to a defined thickness in a Carver laboratory press. This is particularly useful for applications requiring placing a thin film between anatomical structures where space is limited.

Mechanical testing of the freeze-dried material, 15 rehydrated in normal saline, resulted in force to break values of 170 - 900 g/cm². The elongation to break values for this material were between 33 and 62%.

Example 18: This example illustrates the preparation of CMC-modified HA.

20 HA (0.4% w/w, 0.01M) and Aqualon-type CMC having a molecular weight of 250,000 and a degree of substitution in the range 0.65 to 0.90 (0.19% w/w, 0.01M) were mixed together in aqueous solution at room temperature. The pH of the mixture was adjusted to and maintained at pH 4.7 - 4.8 25 by addition of 1M HCl. To each 100 ml of this solution was added 0.67 g (0.04M) EDC. During reaction with EDC, the pH of the solution was maintained at pH 4.7 - 4.8 by addition of 0.1M HCl and the reaction allowed to proceed for 1 hour, during which time a precipitate formed. The unreacted EDC 30 was removed from the precipitate by dialysis against acidified water (pH 4.0) for 24 hours with 2 dialysate changes at 3 and 19 hours. The HA/CMC slurry was then cast

- 24 -

into flat molds and air dried for 24 hours at room temperature.

HA/CMC membranes were shown to reduce the incidence of postoperative adhesion formation in experimental animal models. In experiments using the rat cecal abrasion model, HA/CMC membranes were placed around surgically abraded rat ceca; previous studies had demonstrated that adhesions readily formed to the ceca of rats which had been abraded in controlled fashion. Cecal adhesions in animal groups that received either HA/CMC membranes or ORC membranes (Interceed TC7 membranes marketed by Johnson & Johnson for adhesion prevention) were compared to adhesion controls in animals whose ceca were abraded but did not receive any membrane. The results of these experiments showed that the HA/CMC membranes consistently reduced adhesion formation compared to control animals and to animals that received the Interceed TC7 film.

Example 19: This example illustrates the preparation of EDC-activated HA.

HA (1.0×10^6 daltons) was dissolved in water to make a 0.8% w/v solution by stirring overnight at 25°C. The pH of the reaction mixture was adjusted to pH 4.75 with 0.1N HCl. EDC (4:1 molar ratio of EDC to HA, 1.53% w/v final concentration) was added to this solution with continuous stirring and was maintained at a constant pH (4.7 - 5.1) for one hour by adding additional 0.1N HCl. Removal of the unreacted EDC and other low molecular weight impurities was performed by either molecular weight sizing, dialysis, or diafiltration using standard methods. A water-insoluble, clear gel was obtained after this process.

- 25 -

Example 20: This example illustrates the effect of fractional precipitation of EDC-activated HA with a water soluble solvent.

The procedure described in example 19 was repeated 5 with the exception that unreacted EDC and other low molecular weight impurities were removed by fractional precipitation using a suitable water-soluble solvent (e.g., C₁-C₃ alcohols, acetone). Under these conditions, water insoluble fibers were produced.

10 Example 21: This example illustrates the preparation of EDC-activated CMC.

CMC (250 x 10³ daltons) was dissolved in water to make a 0.8% w/v solution by stirring at room ambient temperature (22-25°C) overnight. The pH of the reaction 15 mixture was adjusted to pH 4.75 with 0.1N HCl. EDC (4:1 molar ratio of EDC to CMC, 1.53% w/v final concentration) was added to this solution with constant stirring and the pH was maintained between 4.70 and 5.10 for one hour by adding additional 0.1N HCl. Removal of the unreacted EDC and other 20 low molecular weight impurities was performed by either molecular weight seizing chromatography, dialysis, diafiltration, or fractional precipitation of the CMC with a suitable water-soluble solvent (e.g., C₁-C₃ alcohols, acetone). Water insoluble fibers, approximately 300 - 800 25 μm long and 10 - 20 μm wide, are produced from these reaction conditions.

Example 22: This example illustrates the preparation of a blend of EDC-activated HA with EDC-activated CMC.

30 EDC-activated HA and CMC were prepared separately as described in Examples 19 and 21 but each reaction product was not purified prior to blending. Three hundred ml of the

- 26 -

activated HA and 300 ml of the activated CMC were placed in a 1000 ml beaker and blended with a Turrax brand blender at 6000 rpm for 10 minutes at 25°C. This resulted mixture was purified by dialysis against pH 4.0 water for 24 hours at a 5 20:1 ratio with 3 dialysate exchanges. After dialysis the mixture was poured into a flat mold and air dried to a thin water insoluble film. The quantity of fibers in the mixture can be controlled by varying the relative amount of activated CMC and activated HA that are blended together.

10 Example 23: This example illustrates the coupling of HA with histidine using HOBr and EDC.

To a solution of sodium hyaluronate (200 mg, 0.5 mmoles, MW 1,700,000) and L-histidine (155.2 mg, 1.0 mmole) in water (40mL) was added 1-hydroxybenzotriazole hydrate (HOBr) (67.6 mg, 0.5 mmoles) followed by adjustment of the pH to 3.35 with 1N HCl. After stirring for one hour, 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (EDC) (154.4 mg, 0.8 mmoles) was added and the pH was maintained between 4.0 and 4.8, by the addition of 1N HCl, 15 for two hours. Saturated sodium carbonate was added to adjust the pH to 7.01 and the product was precipitated by the addition of 95% ethanol (\approx 150 mL). The resulting solid was collected by vacuum filtration, washed with absolute ethanol (3 x 20 mL), and dried by lyophilization. The 20 overall yield of modified polymer was 65%. The material was used directly without further purification.

Example 24: This example illustrates the coupling of HA with 3-dimethylaminopropylamine using HOBr and EDC.

3-Dimethylaminopropylamine (102 mg, 1.0 mmoles) in 30 water (3 mL) was adjusted to pH 7.0 with 1 N HCl and added to a solution of sodium hyaluronate (200 mg, 0.5 mmoles, MW 1,700,000) in water (40 mL). 1-Hydroxybenzotriazole hydrate (HOBr) (67.6 mg, 0.5 mmoles) was added and the pH

- 27 -

adjusted to 3.35 with 1N HCl. After stirring for one hour, 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (EDC) (154.4 mg, 0.8 mmoles) was added and the pH maintained at 4.6 for 50 min. by the addition of 1N HCl and/or 10% w/v 5 sodium carbonate. Saturated sodium carbonate was then added to adjust the pH to 7.0 and half of the product was purified by either the ultrafiltration or ethanol precipitation methods listed below.

An aliquot (20mL) of the reaction solution was 10 purified by ultrafiltration (1,000 MWCO) at ambient temperature against three volume exchanges with water. The retentate was concentrated to about 12 mL and used without further purification.

Another aliquot (20 mL) of the reaction solution was 15 added to 10% w/v aqueous NaCl (5 mL) followed by the addition of 95% ethanol (\approx 75 mL). This resulted in the formation of a precipitate which was collected by vacuum filtration, washed with absolute ethanol (3 x 20 mL), and dried by lyophilization. The product was used directly to 20 prepare a hydrogel without further purification.

Example 25: This example illustrates the coupling of HA with dihydroxyphenylamine (*di*-DOPA) using HOBt and EDC.

To a solution of sodium hyaluronate (200 mg, 0.5 25 mmoles, MW 2,100,000) and *d,l*-DOPA (200 mg, 1.0 mmole) in water (40 mL) was added 1-hydroxy-benzotriazole hydrate (HOBt) (67.6 mg, 0.5 mmoles) followed by adjustment of the pH to 4.35 with 1N HCl. After stirring for one hour, 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (EDC) 30 (154.4 mg, 0.8 mmoles) was added and the pH was maintained between 4.0 and 4.8 for two hours by the appropriate addition of either 1N HCl or 10% w/v aqueous sodium carbonate. Saturated sodium carbonate was added to adjust

- 28 -

the pH to 7.0 and the product was precipitated by the addition of 95% ethanol (\approx 150 mL). The resulting solid was collected by vacuum filtration, washed with absolute ethanol (3 x 20 mL), and dried by lyophilization. The material was 5 used directly to prepare the hydrogel without further purification. Care should be taken to limit the gels exposure of the gel to air in order to protect the material from oxidation.

Example 26: This example illustrates the coupling 10 of CMC with 3-dimethylaminopropylamine using HOBt and EDC.

3-Dimethylaminopropylamine (102 mg, 1.0 mmoles) in water (3 mL) was adjusted to pH 7.0 with 1 N HCl and added to a solution of CMC (125 mg, 0.5 mmoles, MW 250,000) in water (20 mL). 1-Hydroxybenzotriazole hydrate (HOBt) (67.6 15 mg, 0.5 mmoles) was added and the pH adjusted to 4.0 with 1 N HCl. After stirring for one hour, 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (EDC) (154.4 mg, 0.8 mmoles) was added and the pH was maintained at 4.7 for one hour by the addition of 1 N HCl and/or 10% 20 sodium carbonate. At the end of the reaction, 10% sodium carbonate was added to adjust the pH to 7.0. The product was obtained as a white solid after ethanol precipitation.

Example 27: This example illustrates the coupling of Bop-activated HA with glycine methyl ester.

25 To a solution of sodium hyaluronate (200 mg, 0.5 mmoles, MW 2,100,000) in water (40 mL) was added benzotriazole-1-yloxytris(dimethylamino)phosphonium hexafluorophosphate (Bop) (442.3 mg, 1.0 mmoles) in dimethylformamide (1 mL) followed by pH adjustment to 4.6 with 1 N 30 HCl. After stirring for 15 min., glycine methyl ester hydrochloride (126 mg, 1.0 mmoles) was added and the reaction mixture was allowed to stir at ambient temperature for 40 hr. The reaction mixture was transferred to a

- 29 -

separatory funnel and extracted with methylene chloride (3 X 50 mL). The aqueous layer was removed and the product was precipitated with 95% ethanol and collected by vacuum filtration. The solid was washed with a small quantity of 5 absolute ethanol and air dried. This material was used to prepare the hydrogel directly without further purification.

Example 28: This example illustrates the coupling of Bop-activated HA with 3-amino-1-propanol.

To a sodium hyaluronate (200 mg, 0.5 mmoles, 10 MW 1,700,000) solution in water (20 mL) was added benzotriazole-1-yloxytris(dimethylamino)phosphonium-hexafluorophosphate (442.3 mg, 1.0 mmoles). After 15 min, 3-amino-1-propanol (45.1 mg, 0.6 mmoles) was added with dimethylformamide (3 mL) and the pH of the reaction mixture 15 was adjusted to 4.8 with 1.0 N HCl. After stirring overnight at ambient temperature, the pH of this reaction mixture was adjusted to \approx 7.0 with saturated sodium carbonate and the reaction mixture was extracted with methylene chloride (3 X 50 mL). The aqueous phase was 20 separated and 95% ethanol (\approx 100 mL) was added to precipitate the product. The solid was collected by vacuum filtration, washed with absolute ethanol (3 X 20 mL), and dried under reduced pressure with a yield of 60%. The material was used directly to prepare the hydrogel without 25 further purification.

Example 29: This example illustrates the formation of a hydrogel using the reactions products of examples 23 through 28.

A solid portion of any of the reaction products of 30 examples 23 through 28 (20 mg) was added to a 0.9% w/v sodium chloride solution (20 mL), and the mixture was allowed to stand overnight at ambient temperature. After this time period, a clear, colorless hydrogel formed which

- 30 -

was isolated by decanting the remaining sodium chloride solution. These resulting gels then were used directly for physical and *in vivo* evaluations.

Example 30: This example illustrates the coupling 5 of HA and CMC with 3-dimethylaminopropylamine using HOBt and EDC.

To a solution of sodium hyaluronate (4.0g, 10 mmoles, MW 2,300,000) and carboxymethyl cellulose (5.1g, 20 mmoles, MW 250,000) in water (500mL) was added 10 1-hydroxybenzotriazole hydrate (4.1g, 30 mmoles) and 3-dimethylaminopropylamine hydrochloride (4.2g, 30 mmoles) followed by pH adjustment to 4.60 with 1 N HCl. After all the chemicals dissolved, 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (8.0g, 40mmoles) was added 15 and the pH maintained between 4.6 and 5.0 by the addition of either 1 N HCl, or 10% Na₂CO₃, for 1 hour. The pH was then adjusted to 5.0 with 10% Na₂CO₃ and half of the solution was ethanol precipitated with 4X the reaction volume of ethanol, washed with additional ethanol, and air dried. The 20 remaining half was dialyzed (12,000 NMW cutoff dialysis tubing) against water adjusted to pH 4.75 (20X volumes) for 24 hrs.

Use

The films, foams, or gels of the invention can be 25 used as a surgical aid, to prevent adhesions or accretions of body tissues during a post-operation or healing period, following procedures known in the surgical arts, as described, for example, in DeBelder et al., PCT Publication No. WO 86/00912. During surgery one or more pieces of the 30 gel or film, as appropriate, are inserted or injected between or among the tissues that are to be kept separate.

- 31 -

The insoluble materials of the invention can also be used as surface pacification agents, both covalently and/or non-covalently attached to biodurable and erodible polymer surfaces; as sealing agents in anastomotic sites for 5 catheters, bowel anastomosis, endoscopic surgical procedure, vascular grafts, and any prosthetic device requiring gluing together or sealing of potential leakage sites; as a potentially new biocompatible fiber for processing into thread, braids, woven and non-woven webs, weaves, and mats, 10 and sutures for wound closure; sclerosing agents for varicose vein removal, tumors, and aneurism; artificial extracellular matrix material for tissue replacement in skin lacerations and burns.

Films, foams or gels of the invention can further be 15 used for sustained release drug delivery. The drug to be delivered can be covalently bonded to the gel or film, as described, for example, in R.V. Sparer et al., 1983, Chapter 6, pages 107-119, in T.J. Roseman et al., Controlled Release Delivery Systems, Marcel Dekker, Inc., New York; and the gel 20 or film can then be implanted or injected at the locus where delivery is desired.

Other Embodiments

Other embodiments are within the following claims. For example, the scale of the reactions may be increased for 25 commercial production of the compositions of the invention. It will also be well understood by those skilled in the art that varying the ratio of polyanionic polysaccharide to activating agent will control the degree of functionalization of the polyanionic polysaccharide.

- 32 -

Claims

1 1. A method for making a water insoluble
2 biocompatible composition, said method comprising combining,
3 in an aqueous mixture, a polyanionic polysaccharide, a
4 nucleophile, and an activating agent under conditions
5 sufficient to form said composition.

1 2. The method of claim 1 wherein two or more
2 polyanionic polysaccharides are employed.

1 3. The method of claim 1 or 2 wherein said
2 polyanionic polysaccharides are chosen from the group
3 consisting of carboxymethyl cellulose, carboxymethyl
4 amylose, hyaluronic acid, chondroitin-6-sulfate, dermatan
5 sulfate, heparin, and heparin sulfate.

1 4. The method of claim 1 wherein said polyanionic
2 polysaccharide is hyaluronic acid.

1 5. The method of claim 1 wherein said polyanionic
2 polysaccharide is carboxymethyl cellulose.

1 6. The method of claim 1 wherein said polyanionic
2 polysaccharide is carboxymethyl amylose.

1 7. The method of claim 2 wherein two of said
2 polyanionic polysaccharides are hyaluronic acid and
3 carboxymethyl cellulose.

1 8. The method of claim 1 wherein said activating
2 agent is chosen from the group consisting of benzotriazole-
3 1-yloxytris(dimethylamino)phosphonium hexafluorophosphate,
4 O-benzotriazole-1-yl-N,N,N',N'-tetramethyluronium

- 33 -

5 hexafluorophosphate, bromotris(dimethylamino)phosphonium
6 hexafluorophosphate, bromotris(pyrrolidinyl)phosphonium
7 hexafluorophosphate and the corresponding halide salts
8 thereof.

1 9. The method of claim 1 wherein said polyanionic
2 polysaccharide are present in a concentration of 0.0002 -
3 0.1M.

1 10. The method of claim 9 wherein said polyanionic
2 polysaccharide is present in a concentration of 0.0005 -
3 0.02M.

1 11. The method of claim 1 wherein said method is
2 carried out at a pH 3.5 - 8.0.

1 12. The method of claim 1 wherein the stoichiometry
2 of said activating agent to said polysaccharide is at least
3 0.1 molar equivalent of said activating agent per molar
4 equivalent of said polyanionic polysaccharide.

1 13. The method of claim 1 wherein said nucleophile
2 is chosen from the group consisting of an amino acid amide,
3 a monofunctional amine, an amino acid ester, an amino
4 alcohol, an amino thiol, an amino phenol, an amino catechol,
5 an amino acid, a salt of an amino acid, a peptide, and a
6 protein.

1 14. The method of claim 1 wherein the stoichiometry
2 of said polyanionic polysaccharide to said nucleophile is at
3 least 1 molar equivalent of nucleophile per molar equivalent
4 of polyanionic polysaccharide.

- 34 -

1 15. A method for making a water insoluble
2 biocompatible composition, said method comprising combining,
3 in an aqueous mixture, one or more polyanionic
4 polysaccharides, a modifying compound, a nucleophile, and an
5 activating agent under conditions sufficient to form said
6 composition wherein said modifying compound causes the
7 formation of a new active carbonyl groups on said
8 polyanionic polysaccharide.

1 16. The method of claim 15 wherein two or more
2 polyanionic polysaccharides are employed.

1 17. The method of claim 15 or 16 wherein said
2 polyanionic polysaccharides are chosen from the group
3 consisting of carboxymethyl cellulose, carboxymethyl
4 amylose, hyaluronic acid, chondroitin-6-sulfate, dermatan
5 sulfate, heparin, and heparin sulfate.

1 18. The method of claim 15 wherein said polyanionic
2 polysaccharide is hyaluronic acid.

1 19. The method of claim 15 wherein said polyanionic
2 polysaccharide is carboxymethyl cellulose.

1 20. The method of claim 15 wherein said polyanionic
2 polysaccharide is carboxymethyl amylose.

1 21. The method of claim 16 wherein two of said
2 polyanionic polysaccharides are hyaluronic acid and carboxyl
3 methyl cellulose.

1 22. The method of claim 15 wherein said modifying
2 compound is chosen from the group consisting of

- 35 -

3 1-hydroxybenzotriazole hydrate, 1-hydroxybenzotriazole
4 monohydrate, N-hydroxysulfosuccinimide,
5 N-hydroxysuccinimide, 4-nitrophenol, 2-nitrophenol,
6 4-nitrothiophenol, 2-nitrothiophenol, pentachlorophenol,
7 pentafluorophenol, imidazole, tetrazole, and
8 4-dimethylaminopyridine.

1 23. The method of claim 15 wherein said activating
2 agent comprises a carbodiimide.

1 24. The method of claim 23 wherein said
2 carbodiimide comprises 1-ethyl-3-(3-dimethylaminopropyl)
3 carbodiimide, or 1-ethyl-3-(3-dimethylaminopropyl)
4 carbodiimide methiodide.

1 25. The method of claim 15 wherein said polyanionic
2 polysaccharide is present in a concentration of 0.0002 -
3 0.1M.

1 26. The method of claim 25 wherein said polyanionic
2 polysaccharide is present in a concentration of 0.0005 to
3 0.02M.

1 27. The method of claim 15 wherein said method is
2 carried out at a pH 3.5 - 8.0.

1 28. The method of claim 15 wherein the
2 stoichiometry of said polyanionic polysaccharide to said
3 activating agent is at least 0.1 molar equivalent of said
4 activating agent per molar equivalent of said polyanionic
5 polysaccharide.

- 36 -

1 29. The method of claim 15 wherein the
2 stoichiometry of said modifying agent to said activating
3 agent is at least 1 molar equivalent of said modifying
4 compound per molar equivalent of said activating agent.

1 30. The method of claim 15 wherein said nucleophile
2 is chosen from the group consisting of an amino acid amide,
3 a monofunctional amine, an amino acid ester, an amino
4 alcohol, an amino thiol, an amino phenol, an amino catechol,
5 an amino acid, a salt of an amino acid, a peptide, and a
6 protein.

1 31. A water insoluble composition prepared
2 according to the method of claim 1, 2, 15 or 16.

1 32. The composition of claim 31 wherein said
2 composition is in the form of a gel.

1 33. The composition of claim 31 wherein said
2 composition is in the form of fibers.

1 34. The composition of claim 31 wherein said
2 composition is in the form of a membrane.

1 35. The composition of claim 31 wherein said
2 composition is in the form of a foam.

1 36. The composition of claim 31 wherein said
2 composition is in the form of an adhesion prevention
3 composition.

- 37 -

1 37. The composition of claim 31, further comprising
2 a pharmaceutically active substance dispersed within said
3 composition.

1 38. The composition of claim 37 wherein said
2 pharmaceutically active substance is chosen from the group
3 consisting of proteins, growth factors, enzymes, drugs,
4 biopolymers, and biologically compatible synthetic polymers.

1 39. A water insoluble composition comprising the
2 reaction product of a polyanionic polysaccharide, a
3 nucleophile, and an activating agent.

1 40. A water insoluble composition comprising the
2 reaction product of two or more polyanionic polysaccharides,
3 a nucleophile, and an activating agent.

1 41. The water insoluble composition of claim 39 or
2 40 wherein said activating agent is chosen from the group
3 consisting of benzotriazole-1-yloxytris(dimethylamino)-
4 phosphonium hexafluorophosphate, O-benzotriazole-1-yl-
5 N,N,N',N'-tetramethyluronium hexafluorophosphate,
6 bromotris(dimethylamino)phosphonium hexafluorophosphate,
7 bromotris(pyrrolidinyl)phosphonium hexafluorophosphate and
8 the corresponding halide salts thereof.

1 42. A water insoluble composition comprising the
2 reaction product of a polyanionic polysaccharide, a
3 modifying compound, a nucleophile, and an activating agent.

1 43. A water insoluble composition comprising the
2 reaction or product of two or more polyanionic
3 polysaccharides, a modifying compound, a nucleophile, and an
4 activating agent.

1 44. The composition of claim 39, 40, 42 or 43
2 wherein said polyanionic polysaccharides are chosen from the
3 group consisting of carboxymethyl cellulose, carboxymethyl
4 amylose, hyaluronic acid, chondroitin-6-sulfate, dermatin
5 sulfate, heparin, and heparin sulfate.

1 45. The composition of claim 39 or 42 wherein said
2 polyanionic polysaccharide is hyaluronic acid.

1 46. The composition of claim 39 or 42 wherein said
2 polyanionic polysaccharide is carboxymethyl cellulose.

1 47. The composition of claim 39 or 42 wherein said
2 polyanionic polysaccharide is carboxymethyl amylose.

1 48. The composition of claim 40 or 43 wherein two
2 of said polyanionic polysaccharides are hyaluronic acid and
3 carboxy methyl cellulose.

1 49. The composition of claim 39, 40, 42 or 43
2 wherein said nucleophile is chosen from the group consisting
3 of an amino acid amide, a monofunctional amine, an amino
4 acid ester, an amino alcohol, an amino thiol, an amino
5 phenol, an amino catechol, an amino acid, a salt of an amino
6 acid, a peptide, and a protein.

1 50. The composition of claim 42 or 43 wherein said
2 modifying compound is chosen from the group consisting of
3 1-hydroxybenzotriazole hydrate, 1-hydroxybenzotriazole
4 monohydrate, N-hydroxysulfosuccinimide,
5 N-hydroxysuccinimide, 4-nitrophenol, 2-nitrophenol,

- 39 -

6 4-nitrothiophenol, 2-nitrothiophenol, pentachlorophenol,
7 pentafluorophenol, imidazole, tetrazole, and
8 4-dimethylaminopyridine.

1 51. The composition of claim 42 or 43 wherein said
2 activating agent comprises a carbodiimide.

1 52. The composition of claim 51 wherein said
2 carbodiimide comprises 1-ethyl-3-(3-dimethylaminopropyl)
3 carbodiimide, or 1-ethyl-3-(3-dimethylaminopropyl)
4 carbodiimide methiodide.

1 53. The composition of claims 39, 40, 42 or 43
2 wherein said composition is in the form of a gel.

1 54. The composition of claims 39, 40, 42 or 43
2 wherein said composition is in the form of fibers.

1 55. The composition of claims 39, 40, 42 or 43
2 wherein said composition is in the form of a membrane.

1 56. The composition of claims 39, 40, 42 or 43
2 wherein said composition is in the form of a foam.

1 57. The composition of claims 39, 40, 42 or 43
2 wherein said composition is in the form of an adhesion
3 prevention composition.

1 58. The composition of claims 39, 40, 42 or 43,
2 further comprising a pharmaceutically active substance
3 dispersed within said composition.

- 40 -

1 59. The composition of claim 58 wherein said
2 pharmaceutically active substance is chosen from the group
3 consisting of proteins, growth factors, enzymes, drugs,
4 biopolymers, and biologically compatible synthetic polymers.

1 60. A method for making a water insoluble
2 biocompatible composition, said method comprising combining,
3 in an aqueous solution, a polyanionic polysaccharide, and an
4 activating agent under conditions sufficient to form said
5 composition.

1 61. The method of claim 60 wherein said polyanionic
2 polysaccharide is chosen from the group consisting of
3 carboxymethylcellulose, carboxymethylamylose, hyaluronic
4 acid, chondroitin-6-sulfate, dermatin sulfate, heparin, and
5 heparin sulfate.

1 62. The method of claim 61 wherein said polyanionic
2 polysaccharide is hyaluronic acid.

1 63. The method of claim 61 wherein said polyanionic
2 polysaccharide is carboxymethylcellulose.

1 64. The method of claim 61 wherein said polyanionic
2 polysaccharide is carboxymethylamylose.

1 65. The method of claim 60 wherein said activating
2 agent comprises a carbodiimide.

1 66. The method of claim 65 wherein said
2 carbodiimide comprises 1-ethyl-3-(3-dimethylaminopropyl)
3 carbodiimide, or 1-ethyl-3-(3-dimethylaminopropyl)
4 carbodiimide methiodide.

- 41 -

1 67. The method of claim 60 wherein said polyanionic
2 polysaccharide is present in a concentration of 0.005 -
3 0.1M.

1

1 68. The method of claim 67 wherein said polyanionic
2 polysaccharide is present in a concentration of 0.01 -
3 0.02M.

1 69. The method of claim 60 wherein said method is
2 carried out at a pH 3.5 - 8.0.

1 70. The method of claim 60 wherein the molar ratio
2 of said polyanionic polysaccharide to said activating agent
3 is at least 1:1

1 71. The method of claim 60 wherein the molar ratio
2 of said polyanionic polysaccharide to said activating agent
3 is about 4:1.

1 72. A water insoluble composition prepared
2 according to the method of claim 1.

1 73. The composition of claim 72 wherein said
2 composition is in the form of a gel.

1 74. The composition of claim 72 wherein said
2 composition is in the form of fibers.

1 75. A water insoluble composition comprising the
2 reaction product of a polyanionic polysaccharide and an
3 activating agent.

- 42 -

1 76. The composition of claim 75 wherein said
2 polyanionic polysaccharide is chosen from the group
3 consisting of carboxymethylcellulose, carboxymethylamylose,
4 hyaluronic acid, chondroitin-6-sulfate, dermatin sulfate,
5 heparin, and heparin sulfate.

1 77. The composition of claim 75 wherein said
2 polyanionic polysaccharide is hyaluronic acid.

1 78. The composition of claim 75 wherein said
2 polyanionic polysaccharide is carboxymethylcellulose.

1 79. The composition of claim 75 wherein said
2 polyanionic polysaccharide is carboxymethylamylose.

1 80. The composition of claim 75 wherein said
2 activating agent comprises a carbodiimide.

1 81. The composition of claim 80 wherein said
2 carbodiimide comprises 1-ethyl-3-(3-dimethylaminopropyl)
3 carbodiimide, or 1-ethyl-3-(3-dimethylaminopropyl)
4 carbodiimide methiodide.

1 82. The composition of claim 80, further comprising
2 a pharmaceutically active substance dispersed within said
3 gel.

1 83. The composition of claim 82 wherein said
2 pharmaceutically active substance is chosen from the group
3 consisting of proteins, growth factors, enzymes, drugs,
4 biopolymers, and biologically compatible synthetic polymers.

- 43 -

1 84. The composition of claim 75 wherein said
2 composition is in the form of an adhesion prevention
3 composition.

1 85. The composition of claim 75 wherein said
2 composition is in the form of a membrane.

1 86. The composition of claim 75 wherein said
2 composition is in the form of a foam.

1 87. A water insoluble mixture comprising a reaction
2 product of a first polyanionic polysaccharide and an
3 activating agent, in admixture with a reaction product of a
4 second polyanionic polysaccharide and an activating agent.

1 88. The mixture of claim 87 wherein said first
2 polyanionic polysaccharide is carboxymethylcellulose and
3 said second polyanionic polysaccharide is hyaluronic acid.

1 89. The mixture of claim 87 wherein said first
2 polyanionic polysaccharide is carboxymethylamylose and said
3 second polyanionic polysaccharide is hyaluronic acid.

1 90. The mixture of claim 87, further comprising a
2 pharmaceutically active substance dispersed within said
3 blend.

1 91. The mixture of claim 87 wherein said
2 pharmaceutically active substance is chosen from the group
3 consisting proteins, of growth factors, enzymes, drugs,
4 biopolymers, and biologically compatible synthetic polymers.

- 44 -

1 92. The mixture of claim 87 wherein said mixture is
2 in the form of an adhesion prevention composition.

1 93. The mixture of claim 87 wherein said mixture is
2 in the form of a membrane.

1 94. The mixture of claim 87 wherein said mixture is
2 in the form of a foam.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US92/04212

A. CLASSIFICATION OF SUBJECT MATTER

IPC(5) :A61K 31/725, 31/715, 31/70; C07H 1/00

US CL :536/21, 22, 56, 55.2, 55.3, 102, 124; 514/53, 54, 56, 57, 60, 62

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 536/21, 22, 56, 55.2, 55.3, 102, 124; 514/53, 54, 56, 57, 60, 62

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US, A, 3,998,974 (ZAFFARONI) 21 DECEMBER 1976, especially column 14, lines 14 and 15, and column 16, lines 30-60.	15-38, 42, 44-47, 49-68, 71-74 and 76-94
Y	US, A, 4,526,714 (FEIJEN ET AL) 02 JULY 1985, entire document.	1-14, 31-39, 41, 44-47, 49, 53-59 and 60-94
Y	US, A, 4,582,865 (BALAZS ET AL) 15 APRIL 1986, column 1, lines 37 and 38.	32, 73 and 82
Y	US, A, 4,810,748 (LARM) 07 MARCH 1989, column 1, lines 29-42.	3, 17, 44, 61 and 76
Y	CARBOHYDRATE RESEARCH, Volume 16, issued 1971, I. Danishefsky et al, "Conversion of Carboxyl Groups of Mucopolysaccharides into Amides of Amino Acid Esters", pages 199-205, entire document.	1-14, 31-43, 41, 44-49, 53-68, 71-74 and 76-94

Further documents are listed in the continuation of Box C. See patent family annex.

* Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
*'A' document defining the general state of the art which is not considered to be part of particular relevance		
*'E' earlier document published on or after the international filing date	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
*'L' document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
*'O' document referring to an oral disclosure, use, exhibition or other means	"&"	document member of the same patent family
*'P' document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search	Date of mailing of the international search report
27 July 1992	17 AUG 1992
Name and mailing address of the ISA/Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231	Authorized officer EVERETT WHITE Telephone No. (703) 308-0196
Facsimile No. NOT APPLICABLE	